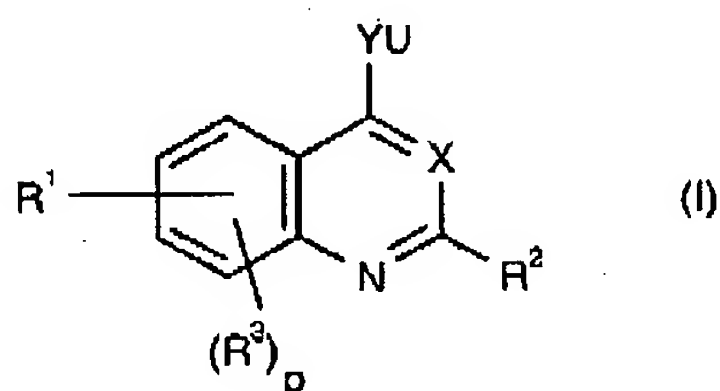


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In the Claims:

A complete listing of claims 1-19 with status identifier follows.

Claim 1 (Currently Amended) A method of treating a susceptible cancer in a human or animal subject in need thereof, comprising administering to said subject an effective amount of a compound of formula (I):



or a salt or solvate thereof;

wherein X is N or CH;

Y is a group W(CH₂), (CH₂)W, or W, in which W is O, S(O)_m wherein m is 0, 1 or 2, or NR^a wherein R^a is hydrogen or a C₁₋₈ alkyl group;

R¹ ~~is represents~~ represents a 5- or 6-membered heterocyclic ring containing 1 to 4 heteroatoms selected from the group N, O or S(O)_m, wherein m is as defined above, with the provisos that the ring does not have two adjacent O or S(O)_m atoms and that where the ring has only N as heteroatom(s) the ring is C-linked to the quinazoline or quinoline ring, R¹ being optionally substituted by one or more R³ groups;

each R³ is independently selected from the group consisting of amino, hydrogen, halogen, hydroxy, nitro, carboxy, formyl, cyano, trifluoromethyl,

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trifluoromethoxy, carbamoyl, ureido, guanidino, C₁₋₈ alkyl, C₁₋₈ alkoxy, C₃₋₈ cycloalkoxyl, C₄₋₈ alkylcycloalkoxy, C₁₋₈ alkylcarbonyl, C₁₋₈ alkoxy carbonyl, N-C₁₋₄ alkylcarbamoyl, N,N-di-[C₁₋₄ alkyl]carbamoyl, hydroxyamino, C₁₋₄ alkoxyamino, C₂₋₄ alkanoyloxyamino, C₁₋₄ alkylamino, di[C₁₋₄ alkyl]amino, di-[C₁₋₄ alkyl]amino-C₁₋₄ alkylene-(C₁₋₄ alkyl)amino, C₁₋₄ alkylamino-C₁₋₄ alkylene-(C₁₋₄ alkyl)amino, hydroxy-C₁₋₄ alkylene-(C₁₋₄ alkyl)amino, phenyl, phenoxy, 4-pyridon-1-yl, pyrrolidin-1-yl, imidazol-1-yl, piperidino, morpholino, thiomorpholino, thiomorpholino-1-oxide, thiomorpholino-1,1-dioxide, piperazin-1-yl, 4-C₁₋₄ alkylpiperazin-1-yl, dioxolanyl, C₁₋₈ alkylthio, arylthio, C₁₋₄ alkylsulphinyl, C₁₋₄ alkylsulphonyl, arylsulphonyl, arylsulphinyl, halogeno-C₁₋₄ alkyl, hydroxy-C₁₋₄ alkyl, C₂₋₄ alkanoyloxy-C₁₋₄ alkyl, C₁₋₄ alkoxy-C₁₋₄ alkyl, carboxy-C₁₋₄ alkyl, formyl-C₁₋₄ alkyl, C₁₋₄ alkoxy carbonyl-C₁₋₄ alkyl, carbamoyl-C₁₋₄ alkyl, N-C₁₋₄ alkylcarbamoyl-C₁₋₄ alkyl, N,N-di-[C₁₋₄ alkyl]carbamoyl-C₁₋₄ alkyl, amino-C₁₋₄ alkyl, C₁₋₄ alkylamino-C₁₋₄ alkyl, di-[C₁₋₄ alkyl]amino-C₁₋₄ alkyl, phenyl-C₁₋₄ alkyl, 4-pyridon-1-yl-C₁₋₄ alkyl, pyrrolidin-1-yl-C₁₋₄ alkyl, imidazol-1-yl-C₁₋₄ alkyl, piperidino-C₁₋₄ alkyl, morpholino-C₁₋₄ alkyl, thiomorpholino-C₁₋₄ alkyl, thiomorpholino-1-oxide-C₁₋₄ alkyl, thiomorpholino-1,1-dioxide-C₁₋₄ alkyl, piperazin-1-yl-C₁₋₄ alkyl, 4-C₁₋₄ alkylpiperazin-1-yl-C₁₋₄ alkyl, hydroxy-C₂₋₄ alkoxy-C₁₋₄ alkyl, C₁₋₄ alkoxy-C₂₋₄ alkoxy-C₁₋₄ alkyl, hydroxy-C₂₋₄ alkylamino-C₁₋₄ alkyl, C₁₋₄ alkoxy-C₂₋₄ alkylamino-C₁₋₄ alkyl, C₁₋₄ alkylthio-C₁₋₄ alkyl, C₁₋₄ alkylsulphinyl-C₁₋₄ alkyl, C₁₋₄ alkylsulphonyl-C₁₋₄ alkyl, hydroxy-C₂₋₄ alkylthio-C₁₋₄ alkyl, C₁₋₄ alkoxy-C₂₋₄ alkylthio-C₁₋₄ alkyl, phenoxy-C₁₋₄ alkyl, anilino-C₁₋₄ alkyl, phenylthio-C₁₋₄ alkyl, cyano-C₁₋₄ alkyl, halogeno-C₂₋₄ alkoxy, hydroxy-C₂₋₄ alkoxy, C₂₋₄ alkanoyloxy-C₂₋₄ alkoxy, C₁₋₄ alkoxy-C₂₋₄ alkoxy, carboxy-C₁₋₄ alkoxy, formyl-

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C₁₋₄ alkoxy, C₁₋₄ alkoxycarbonyl-C₁₋₄ alkoxy, carbamoyl-C₁₋₄ alkoxy, N-C₁₋₄ alkylcarbamoyl-C₁₋₄ alkoxy, N,N-di-[C₁₋₄ alkyl]carbamoyl-C₁₋₄ alkoxy, amino-C₂₋₄ alkoxy, C₁₋₄ alkylamino-C₂₋₄ alkoxy, di-[C₁₋₄ alkyl]amino-C₂₋₄ alkoxy, di-[C₁₋₄ alkyl-C₂₋₄ alkoxy]amino-C₂₋₄ alkoxy, C₂₋₄ alkanoyloxy, hydroxy-C₂₋₄ alkanoyloxy, C₁₋₄alkoxy-C₂₋₄ alkanoyloxy, phenyl-C₁₋₄ alkoxy, phenoxy-C₂₋₄ alkoxy, anilino-C₂₋₄ alkoxy, phenylthio-C₂₋₄ alkoxy, 4-pyridon-1-yl-C₂₋₄ alkoxy, piperidino-C₂₋₄ alkoxy, morpholino-C₂₋₄ alkoxy, thiomorpholino-C₂₋₄ alkoxy, thiomorpholino-1-oxide-C₂₋₄ alkoxy, thiomorpholino-1,1-dioxide-C₂₋₄ alkoxy, piperazin-1-yl-C₂₋₄ alkoxy, 4-C₁₋₄ alkylpiperazin-1-yl-C₂₋₄ alkoxy, pyrrolidin-1-yl-C₂₋₄ alkoxy, imidazol-1-yl-C₂₋₄ alkoxy, halogeno-C₂₋₄ alkylamino, hydroxy-C₂₋₄ alkylamino, C₂₋₄ alkanoyloxy-C₂₋₄ alkylamino, C₁₋₄ alkoxy-C₂₋₄ alkylamino, carboxy-C₁₋₄ alkylamino, C₁₋₄ alkoxycarbonyl-C₁₋₄ alkylamino, carbamoyl-C₁₋₄ alkylamino, N-C₁₋₄ alkylcarbamoyl-C₁₋₄ alkylamino, N,N-di-[C₁₋₄ alkyl]carbamoyl-C₁₋₄ alkylamino, amino-C₂₋₄ alkylamino, C₁₋₄ alkylamino-C₂₋₄ alkylamino, di-[C₁₋₄alkyl]amino-C₂₋₄ alkylamino, phenyl-C₁₋₄ alkylamino, phenoxy-C₂₋₄ alkylamino, anilino-C₂₋₄ alkylamino, 4-pyridon-1-yl- C₂₋₄ alkylamino, pyrrolidin-1-yl-C₂₋₄ alkylamino, imidazol-1-yl-C₂₋₄ alkylamino, piperidino-C₂₋₄ alkylamino, morpholino-C₂₋₄ alkylamino, thiomorpholino-C₂₋₄ alkylamino, thiomorpholino-1-oxide-C₂₋₄ alkylamino, thiomorpholino-1,1-dioxide-C₂₋₄ alkylamino, piperazin-1-yl-C₂₋₄alkylamino, 4-(C₁₋₄alkyl)piperazin-1-yl-C₂₋₄alkylamino, phenylthio-C₂₋₄ alkylamino, C₂₋₄ alkanoylamino, C₁₋₄ alkoxycarbonylamino, C₁₋₄ alkylsulphonylamino, C₁₋₄ alkylsulphinylamino, benzamido, benzenesulphonamido, 3-phenylureido, 2-oxopyrrolidin-1-yl, 2,5-dioxopyrrolidin-1-yl, halogeno-C₂₋₄ alkanoylamino, hydroxy-C₂₋₄ alkanoylamino, hydroxy-C₂₋₄ alkanoyl-(C₁₋₄ alkyl)-amino, C₁₋₄ alkoxy-C₂₋₄ alkanoylamino,

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carboxy-C₂₋₄ alkanoylamino, C₁₋₄ alkoxycarbonyl-C₂₋₄ alkanoylamino, carbamoyl-C₂₋₄ alkanoylamino, N-C₁₋₄ alkylcarbamoyl-C₂₋₄ alkanoylamino, N,N-di-[C₁₋₄ alkyl]carbamoyl-C₂₋₄ alkanoylamino, amino-C₂₋₄ alkanoylamino, C₁₋₄ alkylamino-C₂₋₄ alkanoylamino and ~~or~~ di-[C₁₋₄ alkyl]amino-C₂₋₄ alkanoylamino; and wherein said benzamido or benzenesulphonamido substituent or any anilino, phenoxy or phenyl group on a R³ substituent ~~may~~ optionally has ~~have~~ one or two halogeno, C₁₋₄ alkyl or C₁₋₄ alkoxy substituents; and wherein any substituent having a heterocyclic ring ~~may~~ optionally has ~~have~~ one or two halogeno, C₁₋₄ alkyl or C₁₋₄ alkoxy substituents on said ring; and wherein any substituent having a heterocyclic ring ~~may~~ optionally has ~~have~~ one or two oxo or thioxo substituents on said ring;

or R³ is selected from the group consisting of ~~represents a group selected from~~ M¹-M²-M³-M⁴, M¹-M⁵ and ~~or~~ M¹-M²-M^{3'}-M⁶

wherein

M¹ is ~~represents~~ a C₁₋₄ alkyl group, wherein optionally a CH₂ group is replaced by a CO group;

M² is ~~represents~~ NR¹² or CR¹²R¹³, in which R¹² and R¹³ each independently are ~~represent~~ H or C₁₋₄ alkyl;

M³ is ~~represents~~ a C₁₋₄ alkyl group;

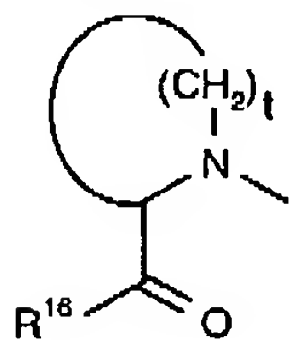
M^{3'} is ~~represents~~ a C₁₋₄ alkyl group or is absent;

M⁴ is selected from the group consisting of ~~represents~~ CN, NR¹²S(O)_mR¹³, S(O)_mNR¹⁴R¹⁵, CONR¹⁴R¹⁵, S(O)_mR¹³ and ~~or~~ CO₂R¹³, in which R¹², R¹³ and m are as defined above and R¹⁴ and R¹⁵ each independently are ~~represent~~ H or C₁₋₄ alkyl, or R¹⁴ and R¹⁵ together with the nitrogen atom to which they are attached form a 5-or 6-membered ring optionally containing 1 or 2 additional heteroatoms selected from N, O or S(O)_m in which ring any nitrogen atom present is ~~may~~

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optionally ~~be~~-substituted with a C₁₋₄ alkyl group, and which ring may optionally ~~has~~ have one or two oxo or thioxo substituents;

~~M⁵ is represents~~ the group NR¹⁴R¹⁵, wherein R¹⁴ and R¹⁵ are as defined above, or ~~M⁵ is represents~~ the group



in which t ~~is represents~~ 2 to 4 and R¹⁶ ~~is represents~~ OH, OC₁₋₄ alkyl or NR¹⁴R¹⁵; and

~~M⁶ is selected from the group consisting of~~ represents a C₃₋₆ cycloalkyl group, the group NR¹⁴R¹⁵, wherein R¹⁴ and R¹⁵ are as defined above, ~~and~~ or a 5- or 6-membered heterocyclic ring system containing 1 to 4 heteroatoms selected from N, O or S;

and p is 0 to 3; or when p is 2 or 3, two adjacent R³ groups together form an optionally substituted methylenedioxy or ethylenedioxy group;

R² is selected from the group consisting of hydrogen, halogen, trifluoromethyl, C₁₋₄ alkyl and C₁₋₄ alkoxy;

U ~~is represents~~ phenyl or a 5 to 10-membered mono or bicyclic ring system in which one or more of the carbon atoms is optionally replaced by a heteroatom independently selected from N, O and S(O)_m, wherein m is 0,1 or 2, and wherein U is substituted by at least one independently selected R⁶ group and U is optionally substituted by at least one independently selected R⁴ group;

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each R^4 is independently selected from the group consisting of hydrogen, hydroxy, halogen, C_{1-4} alkyl, C_{1-4} alkoxy, C_{1-4} alkylamino, di- $[C_{1-4}$ alkyl]amino, C_{1-4} alkylthio, C_{1-4} alkylsulphinyl, C_{1-4} alkylsulphonyl, C_{1-4} alkylcarbonyl, C_{1-4} alkylcarbamoyl, di- $[C_{1-4}$ alkyl] carbamoyl, carbamyl, C_{1-4} alkoxy carbonyl, cyano, nitro and ~~or~~ trifluoromethyl;

each R^6 is independently a group ZR^7 wherein Z is joined to R^7 through a $(CH_2)_p$ group in which p is 0, 1 or 2 and Z is selected from a group consisting of ~~represents a group~~ $V(CH_2)$, $V(CF_2)$, $(CH_2)V$, $(CF_2)V$, $V(CRR')$, $V(CHR)$ and ~~or~~ V where R and R' are each C_{1-4} alkyl and in which V is a hydrocarbonyl group containing 0, 1 or 2 carbon atoms, carbonyl, dicarbonyl, $CH(OH)$, $CH(CN)$, sulphonamide, amide, O, $S(O)_m$ or NR^b where R^b is hydrogen or R^b is C_{1-4} alkyl; and R^7 is an optionally substituted C_{3-6} cycloalkyl; or an optionally substituted 5, 6, 7, 8, 9 or 10-membered carbocyclic or heterocyclic moiety; or R^6 is a group ZR^7 in which Z is NR^b , and NR^b and R^7 together form an optionally substituted 5, 6, 7, 8, 9 or 10-membered carbocyclic or heterocyclic moiety.

Claim 2 (Currently Amended): The A method as claimed in claim 1, wherein the susceptible cancer is a susceptible breast cancer.

Claim 3 (Currently Amended): The A method as claimed in claim 1, wherein the susceptible cancer is a susceptible non-small cell lung cancer.

Claim 4 (Currently Amended): The A method as claimed in claim 1, wherein the susceptible cancer is a susceptible ovarian cancer.

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Claim 5 (Currently Amended): The A method as claimed in claim 1, wherein the susceptible cancer is a susceptible stomach cancer.

Claim 6 (Currently Amended): The A method as claimed in claim 1, wherein the susceptible cancer is a susceptible pancreatic cancer.

Claim 7 (Currently Amended): The A method as claimed in claim 1, wherein the susceptible cancer is a susceptible head and neck cancer.

Claim 8 (Currently Amended): The A method as claimed in claim 1, wherein the susceptible cancer is a susceptible cancer in which there is ~~characterized by~~ expression or over-expression of EGFR.

Claim 9 (Currently Amended): The A method as claimed in claim 1, wherein the susceptible cancer is a susceptible cancer in which there is ~~characterized by~~ expression or over-expression of erbB-2.

Claim 10 (Currently Amended): The A method as claimed in claim 1, wherein the susceptible cancer is a susceptible cancer in which there is ~~characterized by~~ expression or over-expression of EGFR and erbB-2.

Claim 11 (Currently Amended): The A method as claimed in claim 1, wherein X is N.

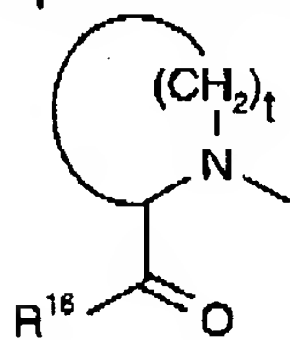
Claim 12 (Currently Amended): The A method as claimed in claim 1, wherein Y is NR^b , $\text{NR}^b(\text{CH}_2)$, or $(\text{CH}_2)\text{NR}^b$, ~~preferably Y is NR^b and R^b is preferably hydrogen or methyl.~~

Claim 13 (Currently Amended): The A method as claimed in claim 1, wherein R^1 is a 5- or 6-membered heterocyclic ring as defined in claim 1 substituted with

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an R^3 group selected from the group consisting of M^1 - M^2 - M^3 - M^4 , M^1 - M^5 and M^1 - M^2 - M^3 - M^6 as defined in claim 1 ~~or claim 2~~; and $p = 0$.

Claim 14 (Currently Amended): The A method as claimed in claim 1, wherein M^1 ~~is~~ represents CH_2 , CO , CH_2CH_2 or CH_2CO ; M^2 ~~is~~ represents NR^{12} in which R^{12} is as defined in claim 1; M^3 ~~is~~ represents CH_2 , CH_2CH_2 or propyl; M^3 ~~is~~ represents CH_2 , ethyl, propyl, isopropyl or is absent; M^4 ~~is~~ represents SOR^{13} , SO_2R^{13} , $NR^{12}SO_2R^{13}$, $SO_2NR^{14}R^{15}$, CO_2R^{13} or $CONR^{14}R^{15}$ in which R^{12} and R^{13} are defined in claim 1 and R^{14} and R^{15} each independently are represent H or C_{1-4} alkyl; M^5 ~~is~~ represents a group $NR^{14}R^{15}$ in which R^{14} and R^{15} together with the nitrogen atom to which they are attached ~~is~~ represent a 6-membered ring optionally containing an additional heteroatom selected from N or O, in which ring any nitrogen atom present ~~is~~ may optionally be substituted with a C_{1-4} alkyl group; or M^5 ~~is~~ represents a group



in which t ~~is~~ represents 2 or 3 and R^{16} ~~is~~ represents OH , NH_2 , $N(C_{1-4} \text{ alkyl})_2$ or OC_{1-4} alkyl; ~~more preferably~~ R^{16} represents NH_2 or $N(CH_3)_2$; or M^5 ~~is~~ represents a group $NR^{14}R^{15}$ in which R^{14} and R^{15} each independently are represent hydrogen or C_{1-4} alkyl, ~~more preferably~~ hydrogen, methyl, ethyl or isopropyl; and M^6 ~~is~~ represents a group $NR^{14}R^{15}$ in which R^{14} and R^{15} each independently ~~is~~ represents C_{1-4} alkyl, ~~more preferably~~ methyl, or R^{14} and R^{15} together with the nitrogen atom to which they are attached ~~is~~ represent a 5- or 6-membered ring optionally containing an additional heteroatom selected from N or O, in which ring any nitrogen atom present ~~is~~ may optionally be substituted with a C_{1-4} alkyl group, ~~preferably a methyl group~~; or M^6 ~~is~~ represents a 5- or 6-membered heterocyclic ring system containing 1 or 2 heteroatoms selected from N or O.

Claim 15 (Currently Amended): The A method as claimed in claim 1, wherein M^2 - M^3 - M^4 ~~is~~ represents a methylsulphonylethylamino, methylsulphinylethylamino, methylsulphonylethyl(methylamino), methylsulphinylethyl(methylamino), methylsulphonylpropylamino, methylsulphinylpropylamino,

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methylsulphonamidoethylamino, aminosulphonylethylamino, methylaminosulphonylethylamino, sarcosinamide, glycine, glycinamide, glycine methyl ester or acetylaminoethylamino group.

Claim 16 (Currently Amended): The A method as claimed in claim 1, wherein R¹ is selected from the group consisting of comprising furan, dihydrofuran, thiophene, imidazole, tetrazole, triazole, pyridine, pyrrole, pyrimidine, isoxazole and or oxadiazole.

Claim 17 (Currently Amended): The A method as claimed in claim 1, wherein R¹ is selected from the group consisting of comprising furan, imidazole, oxadiazole (~~particularly 1,3,4-oxadiazole and 1,2,4-oxadiazole~~) and triazole (~~particularly 1,2,3-triazole and 1,3,4-triazole~~).

Claim 18 (Currently Amended): The A method as claimed in claim 1, wherein R⁶ is benzyl, fluorobenzyl, difluorobenzyl, benzyloxy, fluorobenzyloxy, pyridylmethyl, phenyl, benzenesulphonyl, phenoxy or fluorophenoxy.

Claim 19 (Currently Amended): The A method as claimed in claim 1, wherein U is represents an phenyl, indolyl, isoindolyl, indolinyl, isoindolinyl, 1H-indazolyl, 2,3-dihydro-1H-indazolyl, 1H-benzimidazolyl, 2,3-dihydro-1H-benzimidazolyl or 1H-benzotriazolyl group.

Claim 20 (Currently Amended): The A method as claimed in claim 1, wherein U is represents a phenyl or 1H-indazolyl group.

Claim 21 (Currently Amended): The A method as claimed in claim 1, wherein the optional substituents for the carbocyclic or heterocyclic moiety ~~and also for other optionally substituted groups~~ include hydroxy, halogen, trifluoromethyl, trifluoromethoxy, nitro, amino, cyano, C₁₋₄ alkoxy, C₁₋₄ alkylthio, C₁₋₄ alkyl carbonyl, carboxylate and C₁₋₄ alkoxy carboxyl.

Claim 22 (Currently Amended): The A method as claimed in claim 1, wherein X is represents N; Y is represents NR^a, wherein R^a is hydrogen or C₁₋₄ alkyl; R¹ is

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represents furan, thiophene, pyrrole, pyridine, pyrimidine, pyrazine, imidazole, oxazole, isoxazole, oxadiazole, tetrazole, triazole, dioxolane or a partially or fully hydrogenated derivative of any of these groups, optionally substituted by one or more R^3 groups selected from halo, trifluoromethyl, C_{1-4} alkyl, carboxy, C_{1-4} alkoxy, carbonyl, formyl, hydroxy- C_{1-4} alkyl, 1,3-dioxolan-2-yl, amino, C_{1-4} alkylamino, di(C_{1-4} alkyl)amino, hydroxy- C_{1-4} alkanoyl-(C_{1-4} alkyl)-amino, C_{1-4} alkylamino- C_{1-4} alkyl or di(C_{1-4} alkyl)amino- C_{1-4} alkyl; p is 0; R^2 is represents hydrogen; R^4 is represents hydrogen, halo or methyl; U is represents phenyl, indolyl, benzimidazolyl or indazolyl, more preferably phenyl or indazolyl; and R^6 is represents phenyl, benzyl, α -methylbenzyl, fluorobenzyl, difluorobenzyl, pyridylmethyl, benzenesulphonyl, phenoxy, fluorophenoxy, benzyloxy or fluorobenzyloxy.

Claim 23 (Currently Amended): The A method as claimed in claim 1, wherein X is represents N; Y is represents NR^a , wherein R^a is hydrogen or C_{1-4} alkyl; R^1 is selected from the group consisting of represents a furan, dihydrofuran, thiophene, pyridine, pyrrole, pyrimidine, isoxazole, triazole, tetrazole, imidazole and or oxadiazole ring, preferably furan, imidazole, oxadiazole and triazole, substituted with an R^3 group selected from the group consisting of C_{1-4} alkyl, C_{1-4} alkylamino- C_{1-4} alkyl, di(C_{1-4} alkyl)amino- C_{1-4} alkyl, formyl, carboxy, C_{1-4} alkoxy, carbonyl, dioxolanyl, trifluoromethyl, methylsulphonylethylaminomethyl, methylsulphonylethylamino-carbonyl, methylsulphonylethyl(methylamino)-methyl, methylsulphonamidoethylamino-methyl, aminosulphonylethylamino-methyl, methylaminosulphonylethylamino-methyl, N,N-dimethylaminoprop-2-ylaminomethyl, N-(2-dimethylaminoethyl)-N-ethylaminomethyl, pyridylaminomethyl, tetrahydrofuranomethylaminomethyl, piperazinylmethyl, methylpiperazinylmethyl, piperidinylmethyl, pyridylmethyl, N-(prolinamido)methyl and or (N,N-dimethyl-prolinamido)methyl; p is 0; R^2 is represents hydrogen; R^4 is represents hydrogen or halo; U is represents phenyl or indazolyl; and R^6 is selected from the group consisting of represents benzyl, fluorobenzyl, difluorobenzyl, pyridylmethyl, benzenesulphonyl, phenoxy, benzyloxy or fluorobenzyloxy.